

1401233

- (21) Application No. 46246/72 (22) Filed 6 Oct. 1972
 (31) Convention Application No. 7114/71 (32) Filed 8 Oct. 1971 in
 (33) Czechoslovakia (CS)
 (44) Complete Specification published 23 July 1975
 (51) INT CL² C08F 20/28 8/36 8/40 C08J 7/14
 (52) Index at acceptance
 C3P 102 8D2B2 8D5 8K8 D8 T2A T2D
 (72) Inventor OTTO WICHTERLE



(54) METHOD FOR THE SURFACE TREATMENT OF
 ARTICLES MADE FROM HYDROXYETHYL METHACRYLATE
 POLYMERS

(71) We, CESKOSLOVENSKA
 AKADEMIE VED, a Czechoslovakian
 corporation of No. 3 Národní, Praha 1,
 Czechoslovakia do hereby declare the inven-
 tion, for which we pray that a patent may
 be granted to us, and the method by which
 it is to be performed, to be particularly de-
 scribed in and by the following statement:—

This invention relates to a method for
 modifying the surfaces of articles formed
 from polymers containing free hydroxyl
 groups. More particularly, this invention re-
 lates to a method for modifying the surfaces of
 articles formed from an at least slightly cross-
 linked polymer or copolymer of hydroxyethyl
 methacrylate so as to increase the hydro-
 philic nature thereof and to the articles there-
 by obtained.

Hydrophilic polymers of the type contain-
 ing hydroxyl groups either in the main or
 side chains thereof have assumed consider-
 able practical importance, particularly in the
 cases where their tendency to undergo swell-
 ing in water has been reduced due to the
 effect of the carbon chains in the molecule to
 the point still permitting the realization of
 the desired mechanical properties. This tech-
 nique for reducing swelling in water has
 been advantageously utilized for example, in
 connection with at least slightly cross-linked
 polymers of hydroxyethyl methacrylate. Such
 polymers even when only slightly cross-linked
 do not swell on exposure to water above
 about 45 volume per cent in water.

For various applications of the aforesaid
 type of polymers, as for instance in their
 use as biomedical materials, an improvement
 in their interaction with the surrounding liv-
 ing tissue would be realized if their hydro-
 philic character could be increased without
 any resultant detriment to their mechanical
 properties. This state can be obtained, hereto-
 for only by the application of a surface layer
 of highly hydrophilic material onto the under-
 lying polymeric structure, both the base and
 surface layers being adapted as nearly as pos-
 sible to the single purpose of the required

interaction with the medium in which the
 article is to be maintained. Thus, the base is
 required to be relatively low-swelling and thus
 of a mechanically tough material and the
 covering layer of a less tough, more readily
 swellable material. However, in order to avoid
 any unfavourable mechanical properties due
 to such covering layers, the same have had
 to be very thin and of course, provision has
 to be made so that they adhere perfectly to
 the base.

The experiments which have been carried
 out up until this time in this connection
 have involved the application of the highly
 hydrophilic layers to the base materials by
 means of soluble linear polymers and par-
 ticularly by means of solutions of linear, non-
 cross-linked, hydroxyethyl methacrylate poly-
 mers. However, the adhesion of the films
 prepared in this manner has proved to be
 very unsatisfactory. Proposals in which it has
 been suggested to chemically graft the hydro-
 philic polymers onto free radicals created
 chemically or by radiation on the surface
 of the polymer article have proved somewhat
 more acceptable. However, this latter pro-
 cedure has not been consistently successful
 and then only when considerable effort and
 expense have been involved in that it is almost
 impossible to prepare the thin layer of linear
 polymer and also the structure of the base is
 damaged with the grafting due to the penetra-
 tion by the new chains through it in the
 polymerization of the strongly hydrophilic
 monomer so that the subsequent swelling does
 not only occur at the surface but to a consider-
 able extent into the interior.

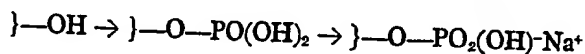
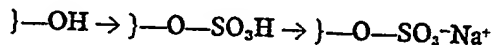
In accordance with the invention it has
 now been found that substantially hydro-
 philic layers can be formed at the surface of
 less hydrophilic at least slightly cross-linked
 polymers and copolymers of hydroxyethyl
 methacrylate, not by the application of an
 additional layer, but by a chemical conversion
 of the surface of the base polymer effected
 only to a relatively small depth.

[Price 33p]

According to the invention there is provided a method for the surface treatment of articles formed from an at least slightly cross-linked polymer or copolymer of hydroxyethyl methacrylate, which method comprises subjecting at least a portion of the surface of such an article to the action of a sulfonation or phosphorylation agent.

Thus in accordance with the invention, it has been found that the formation of a hydro-

philic surface can be achieved by forming new highly hydrophilic groups on the already existing polymer chains which initially contain free hydroxyl groups of the alcoholic type particularly by esterifying such alcoholic groups with sulfuric or phosphoric acid thereby converting the alcohol groups to hydrogen ester groups, which can then be neutralized to form the corresponding alkali or other metallic or onium salts as follows:



The stability of the primary esters of sulfuric or phosphoric acid is entirely adequate even for long term application in aqueous medium, except in the case of the presence of mineral acids (for instance 1 N HCl) where the ester-linked sulfuric and phosphoric acid are rapidly split off.

The above set out substitution i.e. addition reaction can be carried out with any conventional agent used for direct or indirect sulfonation or phosphorylation of alcohol groups. The agents which have proved especially suitable include those having a rate of chemical conversion which is high in relation to the rate of their diffusion into the surface. In such cases, very thin and highly chemically converted surface layers are obtained with only a short exposure of the surface to the action of the agent. Suitable sulfonation agents include 60—100% sulphuric acid, fuming sulphuric acid, gaseous sulphur trioxide alone or mixed with a dry inert gas or gaseous sulphur trioxide dissolved in an organic solvent such as tetrachloro methane or dioxan. Also suitable are chlor-sulphuric and fluoro-sulphuric acids. Use of sulphur trioxide alone results in the formation of a superhydrophilic layer having a thickness of only 1 micron and surprisingly producing a change in the surface properties of the poly (hydroxyethyl methacrylate) hydrophilic gel, without in any way changing its mechanical properties or damaging the microscopic structure of the surface in any way.

Suitable phosphorylation agents include polyphosphoric acid, (especially when prepared from orthophosphoric acid and phosphorus pentoxide) and phosphoryl trichloride.

The process of the invention may be employed advantageously, for example in the superhydrophilization of soft hydrophilic contact lenses. This treatment results in there being obtained a superior sliding i.e., movement of the lens along the cornea. This movement favorably influences the tolerance and non-irritability of the lens. The considerable softness of the superhydrophilic layer is also advantageous in this connection.

If the agent which is selected for use, acts

to swell the material of the base, as for instance, concentrated sulfuric acid or fluoro-sulfuric acid, mixtures of orthophosphoric acid and phosphorus pentoxide or dioxane solutions of sulfur trioxide, deeper layers are formed, for example, amounting to several tenths of a millimeter in thickness. These somewhat thicker layers are suitable for other specific applications because of the continuous transition from the maximally hydrophilized surface layer to the unchanged original material. A substantially perfect adhesion is achieved in this manner, since the internal stresses arising at the transition between layers of different swelling capacity are distributed uniformly throughout the entire volume of the modified layer.

The layers as realized in accordance with the invention differ expressly from a lamination of two different materials as taught by the art in that in the latter, the stresses are concentrated in the proximity of the boundary between the layers.

The procedure according to the invention is particularly adapted for use in the case where improved sliding of the articles is required, i.e., reduced friction as for instance in contact lenses, catheters, surgical sutures and in any application where injury to a living tissue or mucous membrane has to be prevented, packaging and protecting films and foils etc.

Before treatment with the sulphonation or phosphorylation agent, the surface to be treated may be previously impregnated with a compound containing a tertiary amine group, in particular pyridine.

The following examples are further illustrative of this invention, and it will be understood that the invention is not limited thereto.

EXAMPLE 1

A contact lens was manufactured by the conventional mechanical machining of a blank formed by a material comprising a copolymer prepared by polymerization of 12 parts by weight of butyl methacrylate, 87.7 parts by weight of hydroxyethyl methacrylate, and 0.3 parts by weight of ethyl dimethacrylate. Fol-

lowing final polishing to provide the desired optical quality and precise form of the edges, the central portion of the convex area of the lens was covered with paraffin in such a manner, that a border having a width of 1 mm remained uncovered. The lens was then attached by its paraffin covered central portion to a glass rod and, by means of this rod, immersed for 45 seconds into a vessel whose atmosphere was in equilibrium with fuming sulphuric acid containing 60% of sulphur trioxide. After this exposure, the lens was rinsed with water, freed of the paraffin by treatment with toluene, and equilibrated in a 1% solution of sodium bicarbonate. It could then be inserted and used as any other conventionally available contact lenses but it has the advantage of substantially friction and other interruption free movement of the lens area coming into contact with the cornea of the eye. The surface of the front portion of the lens is deliberately not subjected to sulfonation, as an improvement in its sliding movement would make removal of the lens more difficult.

Example 2

The catheter tube was manufactured by a known centrifugal casting method from slightly crosslinked poly(hydroxyethyl methacrylate). The catheter was coated in a dry state on its external surface with a syrupy mixture of polyphosphoric acid, which had been prepared from 3—5 parts of phosphorus pentoxide and 1—2 parts of orthophosphoric acid. The thereby coated catheter was then immersed into a bath of boiling hexane for 3 minutes. The catheter was thereafter rinsed with water, and stored in physiological saline solution diluted with half its volume of 1% aqueous bicarbonate. The catheter can be used directly following the sterilization thereof.

EXAMPLE 3

The exterior of a ship was coated according to known methods using a poly(hydroxyethyl methacrylate) varnish. The coating was then cured with potassium dichromate dissolved in varnish. Thereafter the coating was sulfonated at its surface by moving a vessel supplied with circulating dry air saturated with a mixture of sulfur trioxide vapors corresponding to warm (30°C) fuming sulfuric acid and 60% free SO₃ over the coated area. The exposure of the coating to the vapors amounted to an average of 30 to 60 seconds. As a result the hydrodynamic frictional resistance of the ship was decreased especially after strong swelling of the surface layer.

WHAT WE CLAIM IS:—

1. Method for the surface treatment of articles formed from an at least slightly cross-linked polymer or copolymer of hydroxyethyl methacrylate, which method comprises subjecting at least a portion of the surface of such an article to the action of a sulfonation or phosphorylation agent.
2. Method according to claim 1 wherein the sulfonation agent is 60—100% sulfuric acid.
3. Method according to claim 1 wherein the sulfonation agent is fuming sulfuric acid.
4. Method according to claim 1 wherein the sulfonation agent is gaseous sulfur trioxide.
5. Method according to claim 4 wherein the sulfonation agent is gaseous sulfur trioxide admixed with a dry inert gas.
6. Method according to claim 4 wherein the sulfonation agent is gaseous sulfur trioxide dissolved in an organic solvent.
7. Method according to claim 6 wherein said solvent is tetrachloromethane or dioxan.
8. Method according to claim 1 wherein said sulfonation agent is chlorosulfuric or fluorosulfuric acid.
9. Method according to claim 1 wherein said phosphorylation agent is polyphosphoric acid.
10. Method according to claim 9 wherein said polyphosphoric acid is prepared from orthophosphoric acid and phosphorus pentoxide.
11. Method according to claim 1 wherein said phosphorylation agent is phosphoryl trichloride.
12. Method according to claim 11 wherein the surface of said article has previously been impregnated with a compound containing a tertiary amine group.
13. Method according to claim 12 wherein the surface of said article has previously been impregnated with pyridine.
14. An article having at least a portion of its surface made more hydrophilic than the underlying base portion thereof, produced by the process of claim 1.
15. Method of surface treating articles made of polymers as claimed in claim 1 and substantially as described in any one of the examples herein disclosed.

For the Applicants:
MATTHEWS, HADDAN & CO.,
Chartered Patent Agents,
33 Elmfield Road,
Bromley Kent BR1 1SU.